In the Claims

Claims 1 – 16 (Cancelled)

- 17. (Currently Amended) A molecule of nucleic acid comprising sense and antisense sequences of RNAi placed under the control of a single transcription promoter—of single transcription, the sense and antisense sequences being separated by an intervening a sequence of DNA sequence comprising a transcription stop site sequence for stopping transcription and a gene encoding an antibiotic resistance marker, wherein the intervening DNA sequence is framed at each end thereof by a lox site.
 - 18. (Cancelled)
- 19. (Currently Amended) A pharmaceutical composition comprising a therapeutically effective amount of an active substance of at least one molecule of the nucleic acid of in accordance with claim 17 and a pharmaceutically acceptable compatible excipient.
 - 20. (Cancelled)
- 21. (Currently Amended) A method of <u>transcribing expressing</u> RNAi in cells, comprising:

 a) providing eukaryotic cells and at least one molecule of the nucleic acid of claim 17;
- b) introducing into the eukaryotic cells the at least one molecule of the nucleic acid; a molecule of nucleic acid comprising sense and antisense sequences of RNAi placed under control of a promoter of single transcription, the sense and antisense sequences being separated by a sequence of DNA comprising a sequence for stopping transcription, wherein the DNA sequence is framed at each end thereof by a lox sit, and
- c) providing Cre to the eukaryotic cells such that Cre is placed placing Cre in contact with the lox sites and produces to obtain by site-specific recombination elimination of the intervening DNA.

 sequence and the stop sequence of the transcription such so that the sense and antisense sequences

PHIL1\(\mathbb{3}\)860228.2 2

are only no longer separated except by the [[a]] remaining lox sequences; whereby and thereby permit transcription of the a single RNAi comprising the sense sequence, the lox sequence, and the antisense sequence is transcribed in its entirety with the remaining lox sequence as a loop.

- 22. (Currently Amended) The method according to claim 21, wherein the molecule of the nucleic acid comprises from 5' into 3', a transcription promoter compatible with the <u>eukaryotic</u> cells, the sense sequence of the RNAi, a first lox site, a DNA sequence comprising a <u>transcription stop site</u> and a gene encoding an antibiotic resistance <u>marker-transcription terminator</u>, a second lox site and an antisense sequence of the RNAi.
- 23. (Currently Amended) The method according to claim 21, wherein the molecule of the nucleic acid is a plasmid.
- 24. (Currently Amended) The method according to claim 21, wherein the <u>eukaryotic</u> transfected cells are mammalian cells.
 - 25. (Cancelled).
- 26. (Currently Amended) The method according to claim 21, wherein the <u>antibiotic</u> resistance marker confers resistance to the antibiotic is-neomycin.
- 27. (Currently Amended) The method according to claim 21, wherein <u>Cre is provided to the eukaryotic cells by providing at least one the cells are also transfected with a molecule of a Crc expression nucleic acid comprising a regulating promoter sequence and the cre gene and introducing the Cre expression nucleic acid into the eukaryotic cells.</u>
- 28. (New) The molecule of claim 17 wherein the gene encoding an antibiotic resistance marker confers resistance to the antibiotic neomycin.
- 29. (New) The pharmaceutical composition of claim 19 wherein the gene encoding an antibiotic resistance marker confers resistance to the antibiotic neomycin.

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